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(54) Combinations and compositions which interfere with VEGF/VEGF and angiopoietin/Tie receptor function and their use (II)

(57) The present invention describes the combination of substances interfering with the biological activity of Vascular Endothelial Growth Factor (VEGF)/VEGF receptor systems (compound I) and substances interfering with the biological function of Angiopoietin/Tie receptor systems (compound II) for inhibition of vascularization and for cancer treatment.

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Description

[0001] The present invention provides the combination of substances interfering with the biological activity of Vascular Endothelial Growth Factor (VEGF)/VEGF receptor systems (compound I) and substances interfering with the biological function of Angiopoietin/Tie receptor systems (compound II) for inhibition of vascularization and for cancer treatment.

[0002] Protein ligands and receptor tyrosine kinases that specifically regulate endothelial cell function are substantially involved in physiological as well as in disease-related angiogenesis. These ligand/receptor systems include the Vascular Endothelial Growth Factor (VEGF) and the Angiopoietin (Ang) families, and their receptors, the VEGF receptor family and the tyrosine kinase with immunoglobulin-like and epidermal growth factor homology domains (Tie) family. The members of the two families of receptor tyrosine kinases are expressed primarily on endothelial cells. The VEGF receptor family includes Flt1 (VEGF-R1), Flk1/KDR (VEGF-R2), and Flt4 (VEGF-R3). These receptors are recognized by members of the VEGF-related growth factors in that the ligands of Flt1 are VEGF and placenta growth factor (PlGF), whereas Flk1/KDR binds VEGF, VEGF-C and VEGF-D, and the ligands of Flt4 are VEGF-C and VEGF-D (Nicosia, Am. J. Pathol. 153, 11-16, 1998). The second family of endothelial cell specific receptor tyrosine kinases is represented by Tie1 and Tie2 (also kown as Tek). Whereas Tie1 remains an orphan receptor, three secreted glycoprotein ligands of Tie2, Ang1, Ang2, and Ang3/Ang4 have been discovered (Davis et al., Cell 87, 1161-1169, 1996; Maisonpierre et al., Science 277, 55-60, 1997; Valenzuela et al, Proc. Natl. Acad. Sci. USA 96, 1904-1909, 1999; patents: US 5,521,073; US 5,650,490; US 5,814,464).

[0003] The pivotal role of VEGF and of its receptors during vascular development was exemplified in studies on targeted gene inactivation. Even the heterozygous disruption of the VEGF gene resulted in fatal deficiencies in vascularization (Carmeliet et al., Nature 380, 435-439, 1996; Ferrara et al., Nature 380, 439-442, 1996). Mice carrying homozygous disruptions in either Flt1 or Flk1/KDR gene die in mid-gestation of acute vascular defects. However, the phenotypes are distinct in that Flk1/KDR knock-out mice lack both endothelial cells and a developing hematopoietic system (Shalaby et al. Nature 376, 62-66, 1995), whereas Flt1 deficient mice have normal hematopoietic progenitors and endothelial cells, which fail to assemble into functional vessels (Fong et al., 376, 66-70, 1995). Disruption of the Flt4 gene, whose extensive embryonic expression becomes restricted to lymphatic vessels in adults, revealed an essential role of Flt4 for the remodeling and maturation of the primary vascular networks into larger blood vessels during early development of the cardiovascular system (Dumont et al., Science 282, 946-949, 1998). Consistent with the lymphatic expression of Flt4 in adults overexpression of VEGF-C in the skin of transgenic mice resulted in lymphatic, but not vascular, endothelial proliferation and vessel enlargement (Jeltsch et al., Science 276, 1423-1425, 1997). Moreover, VEGF-C was reported to induce neovascularization in mouse cornea and chicken embryo chorioallantoic membrane models of angiogenesis (Cao et al., Proc. Natl. Acad. Sci. USA 95, 14389-14394, 1998).

[0004] The second class of endothelial cell specific receptor tyrosine kinases has also been found to be critically involved in the formation and integrity of vasculature. Mice deficient in Tie1 die of edema and hemorrhage resulting from poor structural integrity of endothelial cells of the microvasculature (Sato et al., Nature 376, 70-74, 1995; Rodewald & Sato, Oncogene 12, 397-404, 1996). The Tie2 knock-out phenotype is characterized by immature vessels lacking branching networks and lacking periendothelial support cells (Sato et al., Nature 376, 70-74, 1995; Dumont et al., Genes Dev. 8, 1897-1909, 1994). Targeted inactivation of the Tie2 ligand Ang1, as well as overexpression of Ang2, an inhibitory ligand, resulted in phenotypes similar to the Tie2 knock out (Maisonpierre et al., Science 277, 55-60, 1997; Suri et al., cell 87, 1171-1180). Conversely, increased vascularization was observed upon transgenic overexpression of Ang1 (Suri et al., Science 282, 468-471, 1998; Thurstonen et al., Science 286, 2511-2514, 1999).

[0005] The results from angiogenic growth factor expression studies in corpus luteum development (Maisonpierre et al., Science 277, 55-60, 1997; Goede et al. Lab. Invest. 78, 1385-1394, 1998), studies on blood vessel maturation in the retina (Alon et al., Nature Med. 1, 1024-1028, 1995; Benjamin et al, Development 125, 1591-1598, 1998), and gene targeting and transgenic experiments on Tie2, Ang1, and Ang2, suggest a fundamental role of the Angiopoietin/Tie receptor system in mediating interactions between endothelial cells and surrounding pericytes or smooth muscle cells. Ang1, which is expressed by the periendothelial cells and seems to be expressed constitutively in the adult, is thought to stabilize existing mature vessels. Ang2, the natural antagonist of Ang1 which is expressed by endothelial cells at sites of vessel sprouting, seems to mediate loosening of endothelial-periendothelial cell contacts to allow vascular remodeling and sprouting in cooperation with angiogenesis initiators such as VEGF, or vessel regression in the absence of VEGF (Hanahan, Science 277, 48-50, 1997).

[0006] In pathological settings associated with aberrant neovascularization elevated expression of angiogenic growth factors and of their receptors has been observed. Most solid tumors express high levels of VEGF and the VEGF receptors appear predominantly in endothelial cells of vessels surrounding or penetrating the malignant tissue (Plate et al., Cancer Res. 53, 5822-5827, 1993). Interference with the VEGF/VEGF receptor system by means of VEGF-neutralizing antibodies (Kim et al., Nature 362, 841-844, 1993), retroviral expression of dominant negative VEGF receptor variants (Millauer et al., Nature 367, 576-579, 1994), recombinant VEGF-neutralizing receptor variants (Goldman et al., Proc. Natl. Acad. Sci. USA 95, 8795-8800, 1998), or small molecule inhibitors of VEGF receptor tyrosine

kinase (Fong et al., Cancer Res. 59, 99-106, 1999; Wedge et al., Cancer Res. 60, 970-975, 2000; Wood et al. Cancer Res. 60, 2178-2189, 2000), or targeting cytotoxic agents via the VEGF/VEGF receptor system (Arora et al., Cancer Res. 59, 183-188, 1999; EP 0696456A2) resulted in reduced tumor growth and tumor vascularization. However, although many tumors were inhibited by interference with the VEGF/VEGF receptor system, others were unaffected (Millauer et al., Cancer Res. 56, 1615-1620, 1996). Human tumors as well as experimental tumor xenografts contain a large number of immature blood vessels that have not yet recruited periendothelial cells. The fraction of immature vessels is in the range of 40% in slow growing prostate cancer and 90% in fast growing glioblastoma. A selective obliteration of immature tumor vessels was observed upon withdrawal of VEGF by means of downregulation of VEGF transgene expression in a C6 glioblastoma xenograft model. This result is in accordance with a function of VEGF as endothelial cell survival factor. Similarly, in human prostate cancer shutting off VEGF expression as a consequence of androgen-ablation therapy led to selective apoptotic death of endothelial cells in vessels lacking periendothelial cell coverage. In contrast, the fraction of vessels which resisted VEGF withdrawal showed periendothelial cell coverage (Benjamin et al., J. Clin. Invest. 103, 159-165, 1999).

[0007] The observation of elevated expression of Tie receptors in the endothelium of metastatic melanomas (Kaipainen et al., Cancer Res. 54, 6571-6577, 1994), in breast carcinomas (Salvén et al., Br. J. Cancer 74, 69-72, 1996), and in tumor xenografts grown in the presence of dominant-negative VEGF receptors (Millauer et al., Cancer Res. 56, 1615-1620, 1996), as well as elevated expression of Flt4 receptors in the endothelium of lymphatic vessels surrounding lymphomas and breast carcinomas (Jussila et al., Cancer Res. 58, 1599-1604, 1998), and of VEGF-C in various human tumor samples (Salvén et al., Am. J. Pathol. 153, 103-108, 1998), suggested these endothelium-specific growth factors and receptors as candidate alternative pathways driving tumor neovascularization. The high upregulation of Ang2 expression already in early tumors has been interpreted in terms of a host defense mechanism against initial cooption of existing blood vessels by the developing tumor. In the absence of VEGF, the coopted vessels undergo regression leading to necrosis within the center of the tumor. Contrarily, hypoxic upregulation of VEGF expression in cooperation with elevated Ang2 expression rescues and supports tumor vascularization and tumor growth at the tumor margin (Holash et al., Science 284, 1994-1998, 1999; Holash et al., Oncogene 18, 5356-5362, 1999).

[0008] Interference with Tie2 receptor function by means of Angiopoietin-neutralizing Tie2 variants consisting of the extracellular ligand-binding domain has been shown to result in inhibition of growth and vascularization of experimental tumors (Lin et al., J. Clin. Invest. 103, 159-165, 1999; Lin et al. Proc. Natl. Acad. Sci. USA 95, 8829-8834, 1998; Siemeister et al., Cancer Res. 59, 3185-3191, 1999). Comparing the effects of interference with the endothelium-specific receptor tyrosine kinase pathways by means of paracrine expression of the respective extracellular receptor domains on the same cellular background demonstrated inhibition of tumor growth upon blockade of the VEGF receptor system and of the Tie2 receptor system, respectively (Siemeister et al., Cancer Res. 59, 3185-3191, 1999).

It is known that the inhibition of the VEGF/VEGR receptor system by various methods resulted only in slowing down growth of most experimental tumors (Millauer et al., Nature 367, 576-579, 1994; Kim et al., Nature 362, 841-844, 1993; Millauer et al., Cancer Res. 56, 1615-1620, 1996; Goldman et al., Proc. Natl. Acad. Sci. USA 95, 8795-8800, 1998; Fong et al., Cancer Res. 59, 99-106, 1999; Wedge et al., Cancer Res. 60, 970-975, 2000; Wood et al. Cancer Res. 60, 2178-2189, 2000; Siemeister et al., Cancer Res. 59, 3185-3191, 1999). Even by escalation of therapeutic doses a plateau level of therapeutic efficacy was achieved (Kim et al., Nature 362, 841-844, 1993; Wood et al. Cancer Res. 60, 2178-2189, 2000). Similar results were observed upon interference with the Angiopoietin/Tie2 receptor system (Lin et al., J. Clin. Invest. 103, 159-165, 1999; Lin et al., Proc. Natl. Acad. Sci. USA 95, 8829-8834, 1998; Siemeister et al., Cancer Res. 59, 3185-3191, 1999).

[0009] However, there is a high demand for methods that enhance the therapeutic efficacy of anti-angiogenous compounds.

[0010] Searching for methods that enhance the therapeutic efficacy of anti-angiogenic compounds, superior anti-tumor effects were observed unexpectedly upon combination of inhibition of VEGF/VEGF receptor systems and interference with biological function of Angiopoietin/Tie receptor systems. The mode of action underlying the superior effects observed may be that interference biological function of Angiopoietin/Tie receptor systems destabilizes endothelial cell-periendothelial cell interaction of existing mature tumor vessels and thereby sensitizes the endothelium to compounds directed against VEGF/VEGF receptor systems.

[0011] Based on this unexpected finding the present invention provides the combination of functional interference with VEGF/VEGF receptor systems and with Angiopoietin/Tie receptor systems for inhibition of vascularization and of tumor growth.

The pharmaceutical composition consists of two components: compound I inhibits the biological activity of one or several of the VEGF/VEGF receptor systems or consists of cytotoxic agents which are targeted to the endothelium via recognition of VEGF/VEGF receptor systems. Compound II interferes with the biological function of one or several of Angiopoietin/Tie receptor systems or consists of cytotoxic agents which are targeted to the endothelium via recognition of Angiopoietin/Tie receptor systems. Alternatively, compound I inhibits the biological activity of one or several of the VEGF/VEGF receptor systems or of the Angiopoietin/Tie receptor systems and coumpound II consists of cytotoxic

agents which are targeted to the endothelium via recognition of one or several of the VEGF/VEGF receptor systems or of the Angiopoietin/Tie receptor systems. Targeting or modulation of the biological activities of VEGF/VEGF receptor systems and of Angiopietin/Tie receptor systems can be performed by

(a) compounds which inhibit receptor tyrosine kinase activity,
(b) compounds which inhibit ligand binding to receptors,
(c) compounds which inhibit activation of intracellular signal pathways of the receptors,
(d) compounds which inhibit or activate expression of a ligand or of a receptor of the VEGF or Tie receptor system,
(e) delivery systems, such as antibodies, ligands, high-affinity binding oligonucleotides or oligopeptides, or liposomes, which target cytotoxic agents or coagulation-inducing agents to the endothelium via recognition of VEGF/VEGF receptor or Angiopoietin/Tie receptor systems,
(f) delivery systems, such as antibodies, ligands, high-affinity binding oligonucleotides or oligopeptides, or liposomes, which are targeted to the endothelium and induce necrosis or apoptosis.

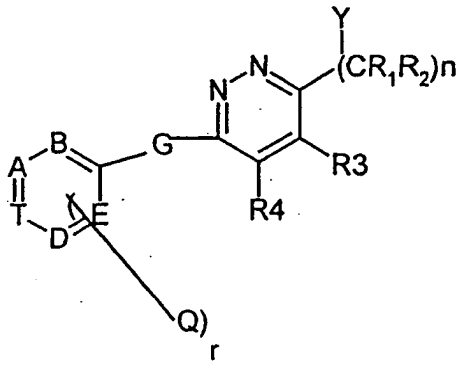
[0012] A compound comprised by compositions of the present invention can be a small molecular weight substance, an oligonucleotide, an oligopeptide, a recombinant protein, an antibody, or conjugates or fusionproteins thereof. An example of an inhibitor is a small molecular weight molecule which inactivates a receptor tyrosine kinase by binding to and occupying the catalytic site such that the biological activity of the receptor is decreased. Kinase inhibitors are known in the art (Sugen: SU5416, SU6668; Fong et al. (1999), Cancer Res. 59, 99-106; Vajkoczy et al., Proc. Am. Associ. Cancer Res. San Francisco (2000), Abstract ID 3612; Zeneca: ZD4190, ZD6474; Wedge et al. (2000), Cancer Res. 60, 970-975; Parke-Davis PD0173073, PD0173074; Johnson et al., Proc. Am. Associ. Cancer Res., San Franzisco (2000), Abstract ID 3614; Dimitroff et al. (1999), Invest. New Drugs 17, 121-135). An example of an antagonist is a recombinant protein or an antibody which binds to a ligand such that activation of the receptor by the ligand is prevented. Another example of an antagonist is an antibody which binds to the receptor such that activation of the receptor is prevented. An example of an expression modulator is an antisense RNA or ribozyme which controls expression of a ligand or a receptor. An example of a targeted cytotoxic agent is a fusion protein of a ligand with a bacterial or plant toxin such as Pseudomonas exotoxin A, Diphtheria toxin, or Ricin A. An example of a targeted coagulation-inducing agent is a conjugate of a single chain antibody and tissue factor. Ligand-binding inhibitors such as neutralizing antibodies which are known in the art are described by Genentech (rhuMAbVEGF) and by Presta et al. (1997), Cancer Res. 57, 4593-4599. Ligand-binding receptor domaines are described by Kendall & Thomas (1993), Proc. Natl. Acad. Sci., U.S.A.90, 10705-10709; by Goldman et al. (1998) Proc. Natl. Acad. Sci., U.S.A.95, 8795-8800 and by Lin et al. (1997), J. Clin. Invest. 100, 2072-2078. Further, dominant negative receptors have been described by Millauer et al. (1994), Nature 367, 567-579.

Receptor blocking antibodies have been described by Imclone (c-p1C11, US 5,874,542). Further known are antagonistic ligand mutants (Slemeister et al. (1998), Proc. Natl. Acad. Sci., U.S.A.95, 4625-4629). High affinity ligand- or receptor binding oligo nucleotides habe been described by NeXstar (NX-244) and Drolet et al. (1996), Nat. Biotech 14, 1021-1025. Further, small molecules and peptides have been described.

[0013] Expression regulators have been described as anti-sense oligo nucleotides and as ribozymes (RPI, Angiozyme™, see RPI Homepage).

[0014] Examples for delivery-/Targeting-Systems have been described as ligand/ antibody-toxin-fusion-proteins or conjugates (Arora et al. (1999), Cancer Res. 59, 183-188 and Olson et al. (1997), Int. J. Cancer 73, 865-870), as endothel cell targeting of liposomes (Spragg et al. (1997), Prog. Natl. Acad. Sci, U.S.A94, 8795-8800, and as endothel cell targeting plus coagulation-induction (Ran et al., (1998), Cancer Res. 58, 4646-4653).

[0015] Small molecules which inhibit the receptor tyrosine kinase activity are for example molecules of general formula I



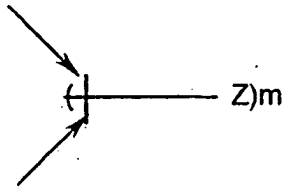
I,

in which

r has the meaning of 0 to 2,

n has the meaning of 0 to 2;

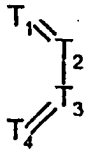
R_3 und R_4 a) each independently from each other have the meaning of lower alkyl,
b) together form a bridge of general partial formula II,



II,

wherein the binding is via the two terminal C- atoms, and

m has the meaning of 0 to 4; or
c) together form a bridge of partial formula III



III,

wherein one or two of the ring members T_1, T_2, T_3, T_4 has the meaning of nitrogen, and each others have the meaning of CH, and the bining is via the atoms T_1 and T_4 ;

G has the meaning of C_1-C_6 - alkyl, C_2 - C_6 - alkylene or C_2 - C_6 - alkenylene; or C_2 - C_6 - alkylene or C_3-C_6- alkenylene, which are substituted with acyloxy or hydroxy; -CH_2-O-, -CH_2-S-, -CH_2-NH-, -CH_2-O-CH_2-, -CH_2-S-CH_2-, -CH_2-NH-CH_2, oxa (-O-), thia (-S-) or imino (-NH-),

A, B, D, E and T independently from each other have the meaning of N or CH, with the provisio that not more than

three of these Substituents have the meaning of N,

Q has the meaning of lower alkyl, lower alkyloxy or halogene,

R_1 and R_2 independently from each other have the meaning of H or lower alkyl,

X has the meaning of imino, oxa or thia;

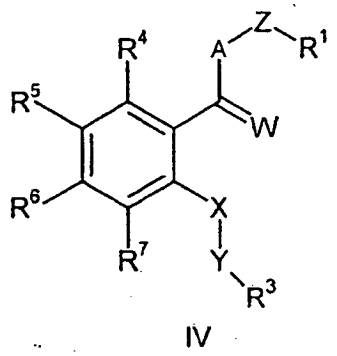
Y has the meaning of hydrogene, unsubstituted or substituted aryl, heteroaryl, or unsubstituted or substituted cycloalkyl; and

Z has the meaning of amino, mono- or disubstituted amino, halogen, alkyl, substituted alkyl, hydroxy, etherificated or esterificated hydroxy, nitro, cyano, carboxy, esterificated carboxy, alkanoyl, carbamoyl, N-mono- or N, N- disubstituted carbamoyl, amidino, guanidino, mercapto, sulfo, phenylthio, phenyl-lower-alkyl-thio, alkyl-phenyl-thio, phenylsulfinyl, phenyl-lower-alkyl-sulfinyl, alkylphenylsulfinyl, phenylsulfonyl, phenyl-lower-alkan-sulfonyl, or alkylphenylsulfonyl, whereas, if more than one rest Z is present ($m \geq 2$), the substituents Z are equal or different from each other, and wherein the bonds marked with an arrow are single or double bonds; or an N-oxide of said compound, wherein one ore more N-atoms carry an oxygene atom, or a salt thereof.

[0016] A preferred salt is the salt of an organic acid, especially a succinate.

[0017] These compounds can preferentially be used as compound I or II in the inventive pharmaceutical composition.

[0018] Compounds which stop a tyrosin phosphorylation, or the persistent angiogenese, respectively, which results in a prevention of tumor growth and tumor spread, are for example anthranyl acid derivatives of general formula IV



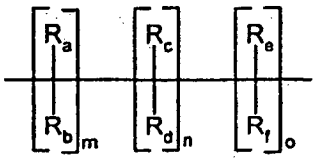
IV

in which

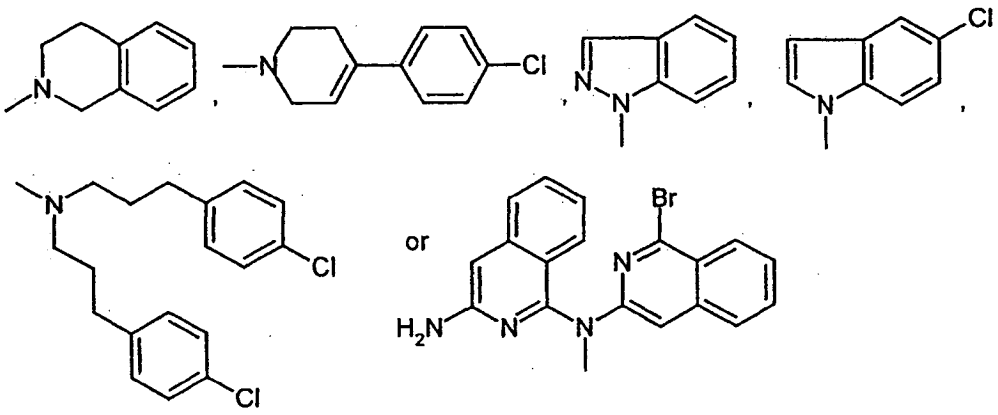
A has the meaning of group $=NR^2$,

W has the meaning of oxygen, sulfur, two hydrogen atoms or the group $=NR^8$,

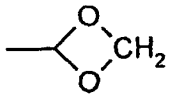
Z has the meaning of the group $=NR^{10}$ or =N-, $-N(R^{10})-(CH_2)_q-$, branched or unbranched C_{1-6}-Alkyl or is the group



or A, Z and R^1 together form the group



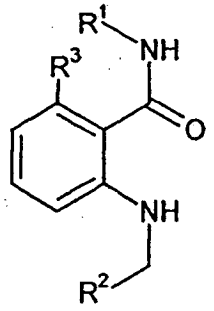
m, n and o	has the meaning of 0 - 3,
q	has the meaning of 1 - 6,
R_a, R_b, R_c, R_d, R_e, R_f	independently from each other have the meaning of hydrogen, C_{1-4} alkyl or the group $=NR^{10}$, and/ or R_a and/ or R_b together with R_c and or R_d or R_c together with R_e and/ or R_f form a bound, or up to two of the groups R_a-R_f form a bridge with each up to 3 C-atoms with R^1 or R^2,
X	has the meaning of group $=NR^9$ or =N-,
Y	has the meaning of group $-(CH_2)_p$,
P	has the meaning of integer 1-4,
R^1	has the meaning of unsubstituted or optionally substituted with one or more of halogene, C_{1-6}-alkyl, or C_{1-6}-alkyl or C_{1-6}-alkoxy, which is optionally substituted by one or more of halogen, or is unsubstituted or substituted aryl or heteroaryl,
R^2	has the meaning of hydrogen or C_{1-6}-alkyl, or form a bridge with up to 3 ring atoms with R_a-R_f together with Z or R_1,
R^3	has the meaning of monocyclic or bicyclic aryl or heteroaryl which is unsubstituted or optionally substituted with one or more of für halogen, C_{1-6}-alkyl, C_{1-6}-alkoxy or hydroxy,
R^4 ,R^5, R^6 and R^7	independently from each other have the meaning of hydrogen, halogen or C_{1-6}-alkoxy, C_{1-6}-alkyl or C_{1-6}-carboxyalkyl, which are unsubstituted or optionally substituted with one or more of halogen, or R^5 and R^6 together form the group



R^8, R^9 and R^{10}	independently from each other have the meaning of hydrogen or C_{1-6}-alkyl, as well as their isomers and salts.

[0019] These compounds can also preferentially be used as compound I or II in the inventive pharmaceutical composition.

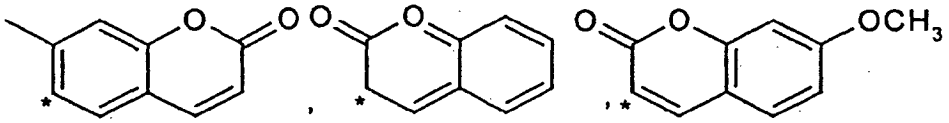
[0020] More preferentially compounds of genearal formula V

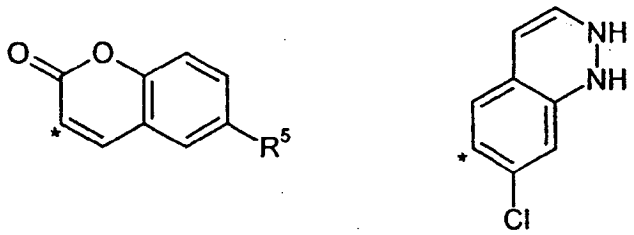


V,

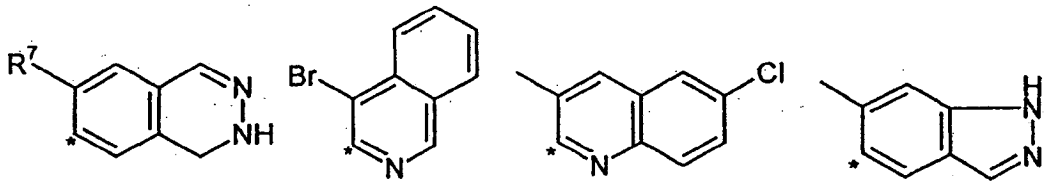
in which

R^1 has the meaning of group

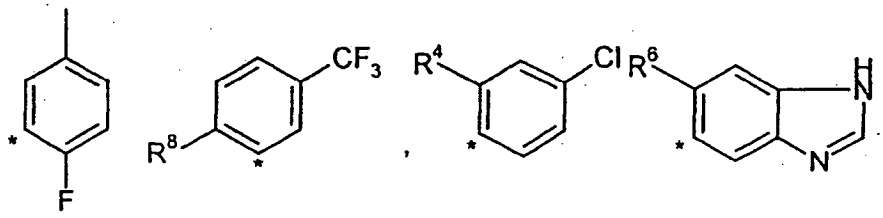




in which R^5 is chloro, bromo or the group $-OCH_3$,



in which R^7 is $-CH_3$ or chloro,

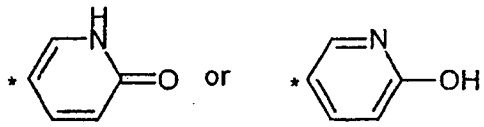


in which R^8 is $-CH_3$, fluoro, chloro or $-CF_3$

in which R^4 is fluoro, chloro, bromo, $-CF_3$, -N=C, $-CH_3$,$-OCF_3$ or $-CH_2OH$

in which R^6 is $-CH_3$ or chloro

R^2 has the meaning of pyridyl or the group



and

R^3 has the meaning of hydrogen or fluoro, as well as their

isomers and salts can be used as compound I or II in the inventive pharmaceutical composition.
These compounds have the same properties as already mentioned above under compound IV and can be used for the treatment of angiogeneous diseases. Compositions comprise compounds of general formulars I, IV and V, alone or in combination.
The above mentioned compounds are also claimed matter within the inventive combinations.

[0021] A further example for ligand binding inhibitors are peptides and DNA sequences coding for such peptides, which are used for the treatment of angiogeneous diseases. Such peptides and DNA sequences are disclosed in Seq. ID No. 1 to 59 of the sequence protocoll. It has been shown that Seq. ID Nos. 34 and 34a are of main interest.

[0022] Claimed matter of the instant invention are therefor pharmaceutical compositions

a) comprising one or several agents as compound I which modulate the biological function of one or several of the VEGF/VEGF receptor systems, and comprising one or several agents as compound II which modulate the biological function of one or several of the Angiopoietin/Tie receptor systems,

b) comprising one or several agents as compound I which are targeted to the endothelium via of one or several of the VEGF/VEGF receptor systems, and comprising one or several agents as compound II which modulate the biological function of one or several of the Angiopoietin/Tie receptor systems,

c) comprising one or several agents as compound I which modulates the biological function of one or several of the VEGF/VEGF receptor systems or of one or several of the Angiopoietin/Tie receptor systems and comprising one or several agents as compound II which are targeted to the endothelium,

d) comprising one or several agents as compound I which modulate the biological function of one or several of the VEGF/VEGF receptor systems, and comprising one or several agents as compound II which are targeted to the endothelium via one or several of the Angiopoietin/Tie receptor systems,

e) comprising one or several agents as compound I which are targeted to the endothelium via one or several of the VEGF/VEGF receptor systems, and comprising one or several agents as compound II which are targeted to the endothelium via one or several of the Angiopoietin/Tie receptor systems,

f) comprising one or several agents as compound I which modulate the biological function of one or several of the VEGF/VEGF receptor systems, and comprising one or several agents as compound II which are targeted to the endothelium via one or several of the VEGF/VEGF receptor systems,

g) comprising one or several agents as compound I which modulate the biological function of one or several of the Angiopoietin/Tie receptor systems, and comprising one or several agents as compound II which are targeted to the endothelium via one or several of the Angiopoietin/Tie receptor systems and

h) comprising one or several agents which interfere with both the function of one or several of the VEGF/VEGF receptor systems and the function of one or several of the Angiopoietin/Tie receptor systems.

[0023] For a sequential therapeutical application the inventive pharmaceutical compositions can be applied simultaneously or separately.

[0024] The inventive compositions comprise as compound I or as compound II at least one of

a) compounds which inhibit receptor tyrosine kinase activity,
b) compounds which inhibit ligand binding to receptors,
c) compounds which inhibit activation of intracellular signal pathways of the receptors,
d) compounds which inhibit or activate expression of a ligand or of a receptor of the VEGF or Tie receptor system,
e) delivery systems, such as antibodies, ligands, high-affinity binding oligonucleotides or oligopeptides, or liposomes, which target cytotoxic agents or coagulation-inducing agents to the endothelium via recognition of VEGF/VEGF receptor or Angiopoietin/Tie receptor systems,
f) delivery systems, such as antibodies, ligands, high-affinity binding oligonucleotides or oligopeptides, or liposomes, which are targeted to the endothelium and induce necrosis or apoptosis.

These compositions are also claimed matter of the present invention.

[0025] Also claimed matter of the present invention are pharmaceutical compositions which comprise as compound I and/ or II at least one of Seq. ID Nos. 1-59. Of most value are pharmaceutical compositions, which comprise as compound I and/ or II Seq. ID Nos. 34a und pharmaceutical compositions according to claims which comprise as compound I and/ or II at least one of sTie2, mAB 4301-42-35, scFv-tTF and/ or L19 scFv-tTF conjugate.

[0026] Further preferred matter of the present invention are pharmaceutical compositions, which comprise as compound I and/ or II at least one small molecule of general formula I, general formula IV and/ or general formula V.

[0027] The most preferred compound which can be used as compound I or II in the inventive composition is (4-Chlorophenyl)[4-(4-pyridylmethyl)-phthalazin-1-yl]ammonium hydrogen succinate.

Therefore, claimed matter of the present invention are also pharmaceutical compositions, which comprise as compound I (4-Chlorophenyl)[4-(4-pyridylmethyl)-phthalazin-1-yl]ammonium hydrogen succinate, sTie2, mAB 4301-42-35, scFv-tTF and/ or L19 scFv-tTF conjugate, and as compound II (4-Chlorophenyl)[4-(4-pyridylmethyl)-phthalazin-1-yl]ammonium hydrogen succinatesTie2, mAB 4301-42-35, scFv-tTF and/ or L19 scFv-tTF conjugate, with the provisio that compound I is not identically to compound II, and most preferred pharmaceutical compositions, which comprise as compound I (4-Chlorophenyl)[4-(4-pyridylmethyl)-phthalazin-1-yl]ammonium hydrogen succinate and as compound II sTie2, mAB 4301-42-35, scFv-tTF and/ or L19 scFv-tTF conjugate; pharmaceutical compositions, which comprise as compound I mAB 4301-42-35 and as compound II sTie2, and/ or scFv-tTF conjugate; pharmaceutical compositions, which comprise as compound I scFv-tTF conjugate and as compound II sTie2 and/ or mAB 4301-42-35; pharmaceutical compositions, which comprise as compound I L19 scFv-tTF conjugate and as compound II sTie2.

[0028] The small molecule compounds, proteins and DNA's expressing proteins, as mentioned above can be used as medicament alone, or in form of formulations for the treatment of tumors, cancers, psoriasis, arthritis, such as rheumatoide arthritis, hemangioma, angiofribroma, eye diseases, such as diabetic retinopathy, neovascular glaukoma, kidney diseases, such as glomerulonephritis, diabetic nephropathy, malignеous nephrosclerose, thrombic microangiopatic syndrome, transplantation rejections and glomerulopathy, fibrotic diseases, such as cirrhotic liver, mesangial cell proliferative diseases, arteriosclerosis and damage of nerve tissues.

The treatment of the damaged nerve tissues with the inventive combination hinders the rapid formation of scars at the damaged position. Thus, there is no scar formation before the axons communicate with each other. Therefore a reconstruction of the nerve bindings is much more easier.

[0029] Further, the inventive combinations can be used for suppression of the ascites formation in patients. It is also possible to suppress VEGF oedemas.

For the use of the inventive combinations as medicament the compounds will be formulated as pharmaceutical composition. Said formulation comprises beside the active compound or compounds acceptable pharmaceutically, organically or inorganically inert carriers, such as water, gelatine, gum arabic, lactose, starch, magnesium stearate, talcum,

plant oils, polyalkylene glycols, etc. Said pharmaceutical preparations can be applied in solid form, such as tablets, pills, suppositories, capsules, or can be applied in fluid form, such as solutions, suspensions or emulsions.
If necessary, the compositions additionally contain additives, such as preservatives, stabilizer, detergents or emulgators, salts for alteration of the osmotic pressure and/ or buffer.
These uses are also claimed matter of the instant invention, as well as the formulations of the active compounds

[0030] For parenteral application especially injectable solutions or suspensions are suitable, especially hydrous solutions of the active compound in polyhydroxyethoxylated castor-oil are suitable.
As carrier also additives can be used, such as salts of the gallic acid or animal or plant phospholipids, as well as mixtures thereof, and liposomes or ingredients thereof.
For oral application especially suitable are tablets, pills or capsules with talcum and/ or hydrocarbon carriers or binders, such as lactose, maize or potato starch.

[0031] The oral application can also be in form of a liquid, such as juice, which optionally contains a sweetener.
The dosis of the active compound differs depending on the application of the compound, age and weight of the patient, as well as the form and the progress of the disease.
The daily dosage of the active compound is 0,5-1000 mg, especially 50-200 mg. The dosis can be applied as single dose or as two or more daily dosis.

[0032] These formulations and application forms are also part of the instant invention.

[0033] Combined functional interference with VEGF/VEGF receptor systems and with Angiopoietin/Tie receptor systems can be performed simultaneously, or in sequential order such that the biological response to interference with one ligand/receptor system overlaps with the biological response to interference with a second ligand/receptor system. Alternatively, combined functional interference with VEGF/VEGF receptor systems or with Angiopoietin/Tie receptor systems and targeting of cytotoxic agents via VEGF/VEGF receptor systems or via Angiopoietin/Tie receptor systems can be performed simultaneously, or in sequential order such that the biological response to functional interference with a ligand/receptor system overlaps in time with targeting of cytotoxic agents.

[0034] The invention is also directed to a substance which functional interferes with both VEGF/VEGF receptor systems and Angiopoietin/Tie receptor systems, or which are targeted via both VEGF/VEGF receptor systems and Angiopoietin/Tie receptor systems.

[0035] VEGF/VEGF receptor systems include the ligands VEGF-A, VEGF-B, VEGF-C, VEGF-D, PlGF, and the receptor tyrosine kinases VEGF-R1 (Flt1), VEGF-R2 (KDR/Flk1), VEGF-R3 (Flt4), and their co-receptors (i.e. neuropilin-1). Angiopoietin/Tie receptor systems include Ang1, Ang2, Ang3/Ang4, and angiopoietin related polypeptides which bind to Tie1 or to Tie2, and the receptor tyrosine kinases Tie1 and Tie2.

[0036] Phamaceutical compositions of the present invention can be used for medicinal purposes. Such diseases are, for example, cancer, cancer metastasis, angiogenesis including retinopathy and psoriasis. Pharmaceutical compositions of the present invention can be applied orally, parenterally, or via gene therapeutic methods.

[0037] Therefor the present invention also concerns the use of pharmaceutical compositions for the production of a medicament for the treatment of tumors, cancers, psoriasis, arthritis, such as rheumatoide arthritis, hemangioma, angiofribroma, eye diseases, such as diabetic retinopathy, neovascular glaukoma, kidney diseases, such as glomerulonephritis, diabetic nephropathie, maligneous nephrosclerosis, thrombic microangiopatic syndrome, transplantation rejections and glomerulopathy, fibrotic diseases, such as cirrhotic liver, mesangial cell proliferative diseases, arteriosclerosis, damage of nerve tissues, suppression of the ascites formation in patients and suppression of VEGF oedemas.

[0038] The following examples demonstrate the feasability of the disclosed invention, without restricting the inventon to the disclosed examples.

Example 1

[0039] Superior effect on inhibition of tumor growth via combination of inhibition of the VEGF A/VEGF receptor system together with functional interference with the Angiopoietin/Tie2 receptor system over separate modes of intervention was demonstrated in an A375v human melanoma xenograft model.

[0040] Human melanoma cell line A375v was stably transfected to overexpress the extracellular ligand-neutralizing domain of human Tie2 receptor tyrosine kinase (sTie2; compound II) (Siemeister et al., Cancer Res. 59, 3185-3191, 1999). For control, A375v cells were stably transfected with the empty expression vector (A375v/pCEP). Swiss *nu/nu* mice were s.c. injected with 1×10^6 transfected A375v/sTie2 or A375v/pCEP tumor cells, respectively. Animals receiving compound I were treated for up to 38 days with daily oral doses of 50 mg/kg of the VEGF receptor tyrosine kinase inhibitor (4-Chlorophenyl)[4-(4-pyridylmethyl)-phthalazin-1-yl]ammonium hydrogen succinate (Wood et al., Cancer Res. 60, 2178-2189, 2000). Various modes of treatment are described in Table 1. Tumor growth was determined by caliper measurement of the largest diameter and its perpendicular.

Table 1

	mode of treatment	
treatment group	(4-Chlorophenyl)[4-(4-pyridylmethyl)-phthalazin-1-yl]ammonium hydrogen succinate (compound I)	sTie2 (compound II)
Group 1: A375v/pCEP	-	-
Group 2: A375v/pCEP	+	-
Group 3: A375v/sTie2	-	+
Group 4: A375v/sTie2	+	+

[0041] Tumors derived from A375v/pCEP control cells reached a size of approx. 250 mm^2 (mean area) within 24 days (Figure 1) without treatment (group 1). Separate treatment with the VEGF receptor inhibitor (4-Chlorophenyl)[4-(4-pyridylmethyl)-phthalazin-1-yl]ammonium hydrogen succinate (compound I, treatment group 2) or separate interference with Angiopoietin/Tie2 receptor system by means of expression of sTie2 (compound II, treatment group 3) delayed growth of tumors to a size of approx. 250 mm^2 to 31 days, respectively. Combination of interference with the Angiopoietin/Tie2 system by means of expression of sTie2 and of interference with the VEGF/VEGF receptor system by means of the kinase inhibitor (4-Chlorophenyl)[4-(4-pyridylmethyl)-phthalazin-1-yl]ammonium hydrogen succinate (compound I + compound II, treatment group 4) delayed growth of the tumors to a size of approx. 250 mm^2 to 38 days. This result clearly demonstrates the superior effect of a combination of interference with the VEGF-A/VEGF receptor system and the Angiopoietin/Tie2 receptor system over separate modes of intervention.

Example 2

[0042] Combination of functional interference with the Angiopoietin/Tie2 receptor system and neutralization of VEGF-A is superior to separate modes of intervention in inhibition of tumor growth.

[0043] Tumors derived from A375v/sTie2 cells and from A375v/pCEP cells were induced in nude mice as described in example 1. Animals receiving compound I were treated twice weekly over a period of time of 4 weeks with intraperitoneal doses of 200 µg of the VEGF-A-neutralizing monoclonal antibody (mAb) 4301-42-35 (Schlaeppi et al., J. Cancer Res. Clin. Oncol. 125, 336-342, 1999). Various modes of treatment are descibed in Table 2. Animals were sacrificed for ethical reasons when tumors of group 1 exceeded a volume of approx. 1000 mm^3. Tumor growth was determined by caliper measurement of the largest diameter and its perpendicular.

Table 2

	mode of treatment	
treatment group	mAb 4301-42-35 (compound I)	sTie2 (compound II)
Group 1: A375v/pCEP	-	-
Group 2: A375v/pCEP	+	-
Group 3: A375v/sTie2	-	+
Group 4: A375v/sTie2	+	+

[0044] Tumors derived from A375v/pCEP control cells reached a size of approx. 1000 mm^3 within 28 days (Figure 2) without treatment (group 1). Tumors treated with the VEGF-A-neutralizing mAb 4301-42-35 (compound I, treatment group 2) grew to a volume of approx. 450 mm^3 within 28 days. Interference with Angiopoietin/Tie2 receptor system by

means of expression of sTie2 (compound II, treatment group 3) reduced growth of tumors within 28 day to a volume of approx. 600 mm^2, respectively. Combination of interference with the Angiopoietin/Tie2 system by means of expression of sTie2 and neutralizing of VEGF-A by means of the mAb 4301-42-35 (compound I + compound II, treatment group 4) resulted in a inhibition of tumor growth to a volume of approx. 250 mm^3 within 28 days.

[0045] The superior effect of a combination of neutralization of VEGF-A and functional interference with the Angiopoietin/Tie2 receptor system over separate modes of intervention is clearly shown.

Example 3

[0046] Combination of functional interference with the Angiopoietin/Tie2 receptor system and targeting of a coagulation-inducing protein via the VEGF/VEGF receptor system is superior to separate modes of intervention in inhibition of tumor growth.

[0047] Tumors derived from A375v/sTie2 cells and from A375v/pCEP cells were induced in nude mice as described in example 1. A single chain antibody (scFv) specifically recognizing the human VEGF-A/VEGF receptor I complex (WO 99/19361) was expressed in E. coli and conjugated to coagulation-inducing recombinant human truncated tissue factor (tTF) by methods descibed by Ran et al. (Cancer Res. 58, 4646-4653, 1998). When tumors reached a size of approx. 200 mm^3 animals receiving compound I were treated on day 0 and on day 4 with intravenous doses of 20 µg of the scFv-tTF conjugate. Various modes of treatment are described in Table 3. Animals were sacrificed for ethical reasons when tumors of group 1 exceeded a volume of approx. 1000 mm^3. Tumor growth was determined by caliper measurement of the largest diameter and its perpendicular.

Table 3

	mode of treatment	
treatment group	scFv-tTF conjugate (compound I)	sTie2 (compound II)
Group 1: A375v/pCEP	-	-
Group 2: A375v/pCEP	+	-
Group 3: A375v/sTie2	-	+
Group 4: A375v/sTie2	+	+

[0048] Tumors derived from A375v/pCEP control cells reached a size of approx. 1000 mm^3 within 28 days (Figure 3) without treatment (group 1). Tumors treated with the coagulation-inducting tTF targeted to the VEGF-A/VEGF receptor I complex via the scFv-tTF conjugate (compound I, treatment group 2) grew to a volume of approx. 500 mm^3 within 28 days. Interference with Angiopoietin/Tie2 receptor system by means of expression of sTie2 (compound II, treatment group 3) reduced growth of tumors within 28 day to a volume of approx. 600 mm^2, respectively. Combination of interference with the Angiopoietin/Tie2 system by means of expression of sTie2 and of targeting the VEGF receptor complex (compound I + compound II, treatment group 4) resulted in a inhibition of tumor growth to a volume of approx. 300 mm^3 within 28 days.

The superior effect of a combination of targeting of the coagulation-inducting tTF to the VEGF-A/VEGF receptor I complex and functional interference with the Angiopoietin/Tie2 receptor system over separate modes of intervention is clearly shown. Similar effects can be expected upon targeting of cytotoxic agents to VEGF/VEGF receptor systems.

Example 4

[0049] Combination of functional interference with the VEGF/VEGF receptor system and targeting of a coagulation-inducing protein via the VEGF/VEGF receptor system is superior to separate modes of intervention in inhibition of tumor growth.

[0050] Tumors derived from A375v/pCEP cells were induced in nude mice as described in example 1. Animals receiving compound I were treated for up to 28 days with daily oral doses of 50 mg/kg of the VEGF receptor tyrosine kinase inhibitor (4-Chlorophenyl)[4-(4-pyridylmethyl)-phthalazin-1-yl]ammonium hydrogen succinate (Wood et al., Cancer Res. 60, 2178-2189, 2000). Compound II consists of a single chain antibody (scFv) specifically recognizing

the human VEGF-A/VEGF receptor I complex (WO 99/19361) which was expressed in E. coli and conjugated to coagulation-inducing recombinant human truncated tissue factor (tTF) by methods descibed by Ran et al. (Cancer Res. 58, 4646-4653, 1998). When tumors reached a size of approx. 200 mm^3 animals receiving compound II were treated on day 0 and on day 4 with intravenous doses of 20 µg of the scFv-tTF conjugate. Various modes of treatment are described in Table 4. Animals were sacrificed for ethical reasons when tumors of group 1 exceeded a volume of approx. 1000 mm^3. Tumor growth was determined by caliper measurement of the largest diameter and its perpendicular.

Table 4

	mode of treatment	
treatment group	(4-Chlorophenyl)[4-(4-pyridylmethyl)-phthal-azin-1-yl]ammonium hydrogen succinate (compound I)	scFv-tTF conjugate (compound II)
Group 1: A375v/pCEP	-	-
Group 2: A375v/pCEP	+	-
Group 3: A375v/pCEP	-	+
Group 4: A375v/pCEP	+	+

[0051] Tumors derived from A375v/pCEP control cells reached a size of approx. 1000 mm^3 within 28 days (Figure 4) without treatment (group 1). Separate treatment with the VEGF receptor inhibitor (4-Chlorophenyl)[4-(4-pyridylmethyl)-phthalazin-1-yl]ammonium hydrogen succinate (compound I, treatment group 2) resulted in a reduction of the tumor volumes to approx. 550 mm^3. Tumors treated with the coagulation-inducting tTF targeted to the VEGF-A/VEGF receptor I complex via the scFv-tTF conjugate (compound II, treatment group 3) grew to a volume of approx. 500 mm^3 within 28 days. Combination of inhibition of VEGF receptor tyrosine kinase by means of (4-Chlorophenyl)[4-(4-pyridylmethyl)-phthalazin-1-yl]ammonium hydrogen succinate and of targeting the VEGF receptor complex (compound I + compound II, treatment group 4) resulted in a inhibition of tumor growth to a volume of approx. 400 mm^3 within 28 days.

[0052] The superior effect of a combination of targeting of the coagulation-inducting tTF to the VEGF-A/VEGF receptor I complex and functional interference with the VEGF/VEGF receptor system over separate modes of intervention is clearly shown. Similar effects can be expected upon targeting of cytotoxic agents to Angiopoietin/Tie receptor systems.

Example 5

[0053] Combination of functional interference with the Angiopoietin/Tie2 receptor system and endothelium-specific targeting of a coagulation-inducing protein is superior to separate modes of intervention in inhibition of tumor growth.

[0054] Tumors derived from A375v/sTie2 cells and from A375v/pCEP cells were induced in nude mice as described in example 1. A fusion protein (L19 scFv-tTF) consisting of L19 single chain antibody specifically recognizing the oncofoetal ED-B domain of fibronectin and the extracellular domain of tissue factor was expressed in E. coli as described by Nilsson et al. (Nat. Med., in press). Further, L19 scFv-tTF data have been represented by D. Neri and F. Nilsson (Meeting "Advances in the application of monoclonal antibodies in clinical oncology", Samos, Greece, 31. May-2. June 2000). When tumors reached a size of approx. 200 mm^3 animals receiving compound I were treated with a single intravenous dose of 20 µg of L19 scFv-tTF in 200 µl saline. Various modes of treatment are described in Table 5. Animals were sacrificed for ethical reasons when tumors of group 1 exceeded a volume of approx. 1000 mm^3. Tumor growth was determined by caliper measurement of the largest diameter and its perpendicular.

Table 5

	mode of treatment	
treatment group	L19 scFv-tTF (compound I)	sTie2 (compound II)
Group 1: A375v/pCEP	-	-
Group 2: A375v/pCEP	+	-
Group 3: A375v/sTie2	-	+
Group 4: A375v/sTie2	+	+

[0055] Tumors derived from A375v/pCEP control cells reached a size of approx. 1000 mm^3 within 28 days (Figure 5) without treatment (group 1). Tumors treated with the coagulation-inducting L19 scFv-tTF (compound I, treatment group 2) grew to a volume of approx. 450 mm^3 within 28 days. Interference with Angiopoietin/Tie2 receptor system by means of expression of sTie2 (compound II, treatment group 3) reduced growth of tumors within 28 day to a volume of approx. 600 mm^2, respectively. Combination of interference with the Angiopoietin/Tie2 system by means of expression of sTie2 and of targeting the endothelium with L19 scFv-tTF (compound I + compound II, treatment group 4) resulted in a inhibition of tumor growth to a volume of approx. 250 mm^3 within 28 days.
The superior effect of a combination of targeting of L19 scFv-tTF to the endothelium and functional interference with the Angiopoletin/Tie2 receptor system over separate modes of intervention is clearly shown.

Example 6

[0056] Combination of functional interference with the VEGF/VEGF receptor system and endothelium-specific targeting of a coagulation-inducing protein is superior to separate modes of intervention in inhibition of tumor growth.
[0057] Tumors derived from A375v/pCEP cells were induced in nude mice as described in example 1. Animals receiving compound I were treated for up to 28 days with daily oral doses of 50 mg/kg of the VEGF receptor tyrosine kinase inhibitor (4-Chlorophenyl)[4-(4-pyridylmethyl)-phthalazin-1-yl]ammonium hydrogen succinate (Wood et al., Cancer Res. 60, 2178-2189, 2000). Compound II consists of L19 scFv-tTF fusion protein as described in example 5. When tumors reached a size of approx. 200 mm^3 animals receiving compound II were treated with a single intravenous dose of 20 µg of L19 scFv-tTF in 200 µl saline. Various modes of treatment are described in Table 6. Animals were sacrificed for ethical reasons when tumors of group 1 exceeded a volume of approx. 1000 mm^3. Tumor growth was determined by caliper measurement of the largest diameter and its perpendicular.

Table 6

	mode of treatment	
treatment group	(4-Chlorophenyl)[4-(4-pyridylmethyl)-phthal-azin-1-yl]ammonium hydrogen succinate (compound I)	L19 scFv-tTF (compound II)
Group 1: A375v/pCEP	-	-
Group 2: A375v/pCEP	+	-
Group 3: A375v/pCEP	-	+
Group 4: A375v/pCEP	+	+

[0058] Tumors derived from A375v/pCEP control cells reached a size of approx. 1000 mm^3 within 28 days (Figure 6) without treatment (group 1). Separate treatment with the VEGF receptor inhibitor (4-Chlorophenyl)[4-(4-pyridylme-

thyl)-phthalazin-1-yl]ammonium hydrogen succinate (compound I, treatment group 2) resulted in a reduction of the tumor volumes to approx. 550 mm^3. Tumors treated with the coagulation-inducting L19 scFv-tTF targeted to the endothelium (compound II, treatment group 3) grew to a volume of approx. 450 mm^3 within 28 days. Combination of inhibition of VEGF receptor tyrosine kinase by means of (4-Chlorophenyl)[4-(4-pyridylmethyl)-phthalazin-1-yl]ammonium hydrogen succinate and of targeting the VEGF receptor complex (compound I + compound II, treatment group 4) resulted in a inhibition of tumor growth to a volume of approx. 200 mm^3 within 28 days.

[0059] The superior effect of a combination of targeting of L19 scFv-tTF to the endothelium and functional interference with the VEGF/VEGF receptor system over separate modes of intervention is clearly shown.

Description of the figures

[0060] Fig. 1 shows the superior effect of combination of interference with VEGF/VEGF receptor system by means of an specific tyrosine kinase inhibitor and with the Angiopoietin/Tie2 receptor system by means of a soluble receptor domain on inhibition of tumor growth (treatment modes of groups 1-4 are given in Table 1). The abbreviations have the following meaning:

mock, con.	=	treatment group 1
mock+VEGF-A	=	treatment group 2
sTIE2-cl13	=	treatment group 3
sTIE2-cl13+VEGF-A	=	treatment group 4

[0061] Fig. 2 shows the superior effect on tumor growth inhibition of combination of VEGF-neutralization and functional interference with Angiopoietin/Tie2 receptor system over separate modes of intervention (treatment modes of groups 1-4 are given in Table 2).

[0062] Fig. 3 shows the superior effect on tumor growth inhibition of combination of targeting of the coagulation-inducing tTF to the VEGF/VEGF receptor I complex via a scFv-tTF conjugate and functional interference with Angiopoietin/Tie2 receptor system over separate modes of intervention (treatment modes of groups 1-4 are given in Table 3).

[0063] Fig. 4 shows the superior effect on tumor growth inhibition of combination of targeting of the coagulation-inducing tTF to the VEGF/VEGF receptor I complex via a scFv-tTF conjugate and functional interference with VEGF/VEGF receptor system by means of the VEGF receptor tyrosine kinase inhibitor (4-Chlorophenyl)[4-(4-pyridylmethyl)-phthalazin-1-yl]ammonium hydrogen succinate over separate modes of intervention (treatment modes of groups 1-4 are given in Table 4).

[0064] Fig. 5 shows the superior effect on tumor growth inhibition of combination of targeting of the coagulation-inducing L19 scFv-tTF fusion protein to the endothelium and functional interference with Angiopoietin/Tie2 receptor system over separate modes of intervention (treatment modes of groups 1-4 are given in Table 5).

[0065] Fig. 6 shows the superior effect on tumor growth inhibition of combination of targeting of the coagulation-inducing L19 scFv-tTF fusion protein to the endothelium and functional interference with VEGF/VEGF receptor system by means of the VEGF receptor tyrosine kinase inhibitor (4-Chlorophenyl)[4-(4-pyridylmethyl)-phthalazin-1-yl]ammonium hydrogen succinate over separate modes of intervention (treatment modes of groups 1-4 are given in Table 6).

Annex to the application documents - subsequently filed sequences listing

[0066]

Sequence Identifier

<110> Schering Aktiengesellschaft

<120> Combinations and compositions which interfere with VEGF/ VEGF and angiopoietin/ Tie receptor function and their use II

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<213> Human

<400> 7

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<400> 8

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<211> 561
<212> DNA
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<400> 9

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<400> 10

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<210> 11
<211> 389
<212> DNA
<213> Human

<400> 11

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<400> 12

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<210> 13
<211> 401
<212> DNA
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<400> 13

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<210> 14

<211> 1002
<212> DNA
<213> Human

<400> 14

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<211> 280
<212> DNA
<213> Human

<400> 15

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<211> 2041
<212> DNA
<213> Human

<400> 16

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<400> 32

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<400> 51

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<400> 52

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<212> DNA
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<400> 29

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<400> 34

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<211> 1036
<212> PRT

<213> homo sapiens

<400> 35

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Arg Ala Leu Val Cys Leu Pro Cys Asp Glu Ser Lys Cys Glu Glu Pro
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Arg Asn Arg Pro Gly Ser Ile Val Gln Gly Val Cys Gly Cys Cys Tyr
    50                  55                  60
Thr Cys Ala Ser Gln Gly Asn Glu Ser Cys Gly Gly Thr Phe Gly Ile
65                  70                  75                  80
Tyr Gly Thr Cys Asp Arg Gly Leu Arg Cys Val Ile Arg Pro Pro Leu
                85                  90                  95
Asn Gly Asp Ser Leu Thr Glu Tyr Glu Ala Gly Val Cys Glu Asp Glu
            100                 105                 110
Asn Trp Thr Asp Asp Gln Leu Leu Gly Phe Lys Pro Cys Asn Glu Asn
        115                 120                 125
Leu Ile Ala Gly Cys Asn Ile Ile Asn Gly Lys Cys Glu Cys Asn Thr
    130                 135                 140
Ile Arg Thr Cys Ser Asn Pro Phe Glu Phe Pro Ser Gln Asp Met Cys
145                 150                 155                 160
Leu Ser Ala Leu Lys Arg Ile Glu Glu Glu Lys Pro Asp Cys Ser Lys
                165                 170                 175
Ala Arg Cys Glu Val Gln Phe Ser Pro Arg Cys Pro Glu Asp Ser Val
            180                 185                 190
Leu Ile Glu Gly Tyr Ala Pro Pro Gly Glu Cys Cys Pro Leu Pro Ser
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Arg Cys Val Cys Asn Pro Ala Gly Cys Leu Arg Lys Val Cys Gln Pro
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Cys Cys Asp Leu Tyr Glu Cys Lys Pro Val Phe Gly Val Asp Cys Arg
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Thr Val Glu Cys Pro Thr Val Gln Gln Thr Ala Cys Pro Pro Asp Ser
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Tyr Glu Thr Gln Val Arg Leu Thr Ala Asp Gly Cys Cys Thr Leu Pro
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Thr Arg Cys Glu Cys Leu Ser Gly Leu Cys Gly Phe Pro Val Cys Glu
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Val Gly Ser Thr Pro Arg Ile Val Ser Arg Gly Asp Gly Thr Pro Gly
305                 310                 315                 320
Lys Cys Cys Asp Val Phe Glu Cys Val Asn Asp Thr Lys Pro Ala Cys
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Val Phe Asn Asn Val Glu Tyr Tyr Asp Gly Asp Met Phe Arg Met Asp
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Asn Cys Arg Phe Cys Arg Cys Gln Gly Gly Val Ala Ile Cys Phe Thr
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Ala Gln Cys Gly Glu Ile Asn Cys Glu Arg Tyr Tyr Val Pro Glu Gly
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Glu Cys Cys Pro Val Cys Glu Asp Pro Val Tyr Pro Phe Asn Asn Pro
385                 390                 395                 400
Ala Gly Cys Tyr Ala Asn Gly Leu Ile Leu Ala His Gly Asp Arg Trp
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Arg Glu Asp Asp Cys Thr Phe Cys Gln Cys Val Asn Gly Glu Arg His
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Cys Val Ala Thr Val Cys Gly Gln Thr Cys Thr Asn Pro Val Lys Val
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Pro Gly Glu Cys Cys Pro Val Cys Glu Glu Pro Thr Ile Ile Thr Val
    450                 455                 460
Asp Pro Pro Ala Cys Gly Glu Leu Ser Asn Cys Thr Leu Thr Arg Lys
465                 470                 475                 480
Asp Cys Ile Asn Gly Phe Lys Arg Asp His Asn Gly Cys Arg Thr Cys
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Gln Cys Ile Asn Thr Gln Glu Leu Cys Ser Glu Arg Lys Gln Gly Cys
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Thr Leu Asn Cys Pro Phe Gly Phe Leu Thr Asp Ala Gln Asn Cys Glu
        515                 520                 525
Ile Cys Glu Cys Arg Pro Arg Pro Lys Lys Cys Arg Pro Ile Ile Cys
    530                 535                 540
Asp Lys Tyr Cys Pro Leu Gly Leu Leu Lys Asn Lys His Gly Cys Asp
545                 550                 555                 560
Ile Cys Arg Cys Lys Lys Cys Pro Glu Leu Ser Cys Ser Lys Ile Cys
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Pro Leu Gly Phe Gln Gln Asp Ser His Gly Cys Leu Ile Cys Lys Cys
            580                 585                 590
Arg Glu Ala Ser Ala Ser Ala Gly Pro Pro Ile Leu Ser Gly Thr Cys
        595                 600                 605
Leu Thr Val Asp Gly His His His Lys Asn Glu Glu Ser Trp His Asp
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Gly Cys Arg Glu Cys Tyr Cys Leu Asn Gly Arg Glu Met Cys Ala Leu
625                 630                 635                 640
Ile Thr Cys Pro Val Pro Ala Cys Gly Asn Pro Thr Ile His Pro Gly
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Gln Cys Cys Pro Ser Cys Ala Asp Asp Phe Val Val Gln Lys Pro Glu
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Leu Ser Thr Pro Ser Ile Cys His Ala Pro Gly Gly Glu Tyr Phe Val
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Glu Gly Glu Thr Trp Asn Ile Asp Ser Cys Thr Gln Cys Thr Cys His
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Ser Gly Arg Val Leu Cys Glu Thr Glu Val Cys Pro Pro Leu Leu Cys
705                 710                 715                 720
Gln Asn Pro Ser Arg Thr Gln Asp Ser Cys Cys Pro Gln Cys Thr Asp
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Gln Pro Phe Arg Pro Ser Leu Ser Arg Asn Asn Ser Val Pro Asn Tyr
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Cys Lys Asn Asp Glu Gly Asp Ile Phe Leu Ala Ala Glu Ser Trp Lys
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Pro Asp Val Cys Thr Ser Cys Ile Cys Ile Asp Ser Val Ile Ser Cys
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Phe Ser Glu Ser Cys Pro Ser Val Ser Cys Glu Arg Pro Val Leu Arg
785                 790                 795                 800
Lys Gly Gln Cys Cys Pro Tyr Cys Ile Lys Asp Thr Ile Pro Lys Lys
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Val Val Cys His Phe Ser Gly Lys Ala Tyr Ala Asp Glu Glu Arg Trp
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Asp Leu Asp Ser Cys Thr His Cys Tyr Cys Leu Gln Gly Gln Thr Leu
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Cys Ser Thr Val Ser Cys Pro Pro Leu Pro Cys Val Glu Pro Ile Asn
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Val Glu Gly Ser Cys Cys Pro Met Cys Pro Glu Met Tyr Val Pro Glu
865                 870                 875                 880
Pro Thr Asn Ile Pro Ile Glu Lys Thr Asn His Arg Gly Glu Val Asp
                885                 890                 895
Leu Glu Val Pro Leu Trp Pro Thr Pro Ser Glu Asn Asp Ile Val His
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Leu Pro Arg Asp Met Gly His Leu Gln Val Asp Tyr Arg Asp Asn Arg
        915                 920                 925
Leu His Pro Ser Glu Asp Ser Ser Leu Asp Ser Ile Ala Ser Val Val
    930                 935                 940
Val Pro Ile Ile Ile Cys Leu Ser Ile Ile Ile Ala Phe Leu Phe Ile
945                 950                 955                 960
Asn Gln Lys Lys Gln Trp Ile Pro Leu Leu Cys Trp Tyr Arg Thr Pro
                965                 970                 975
Thr Lys Pro Ser Ser Leu Asn Asn Gln Leu Val Ser Val Asp Cys Lys
            980                 985                 990
Lys Gly Thr Arg Val Gln Val Asp Ser Ser Gln Arg Met Leu Arg Ile
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<212> DNA
<213> homo sapiens

<400> 36

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<212> DNA
<213> homo sapiens

<400> 37

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<211> 134
<212> DNA
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<400> 38

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<210> 39
<211> 644
<212> DNA
<213> homo sapiens

<400> 39

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<210> 40
<211> 657
<212> DNA
<213> homo sapiens

<400> 40

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<211> 1328
<212> DNA
<213> homo sapiens

<400> 41

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<210> 42
<211> 987
<212> DNA
<213> homo sapiens

<400> 42

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<210> 43
<211> 956
<212> DNA
<213> homo sapiens

<400> 43

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<400> 49

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<400> 52

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Claims

1. Pharmaceutical compositions comprising one or several agents as compound I which modulate the biological function of one or several of the VEGF/VEGF receptor systems, and comprising one or several agents as compound II which modulate the biological function of one or several of the Angiopoietin/Tie receptor systems.

2. Pharmaceutical compositions comprising one or several agents as compound I which are targeted to the endothelium via of one or several of the VEGF/VEGF receptor systems, and comprising one or several agents as compound II which modulate the biological function of one or several of the Angiopoietin/Tie receptor systems.

3. Pharmaceutical compositions comprising one or several agents as compound I which modulates the biological function of one or several of the VEGF/VEGF receptor systems or of one or several of the Angiopoietin/ Tie receptor systems and comprising one or several agents as compound II which are targeted to the endothelium.

4. Pharmaceutical compositions comprising one or several agents as compound I which modulate the biological function of one or several of the VEGF/VEGF receptor systems, and comprising one or several agents as compound II which are targeted to the endothelium via one or several of the Angiopoietin/Tie receptor systems.

5. Pharmaceutical compositions comprising one or several agents as compound I which are targeted to the endothelium via one or several of the VEGF/VEGF receptor systems, and comprising one or several agents as compound II which are targeted to the endothelium via one or several of the Angiopoietin/Tie receptor systems.

6. Pharmaceutical compositions comprising one or several agents as compound I which modulate the biological function of one or several of the VEGF/VEGF receptor systems, and comprising one or several agents as compound II which are targeted to the endothelium via one or several of the VEGF/VEGF receptor systems.

7. Pharmaceutical compositions comprising one or several agents as compound I which modulate the biological function of one or several of the Angiopoietin/Tie receptor systems, and comprising one or several agents as compound II which are targeted to the endothelium via one or several of the Angiopoietin/Tie receptor systems.

8. Pharmaceutical compositions comprising one or several agents which interfere with both the function of one or several of the VEGF/VEGF receptor systems and the function of one or several of the Angiopoietin/Tie receptor systems.

9. Pharmaceutical compositions according to claims 1-8 which are intended for simultaneous or separate sequential therapeutical application.

10. Pharmaceutical compositions according to claims 1-8 which comprise as compound I at least one of

a) compounds which inhibit receptor tyrosine kinase activity,
b) compounds which inhibit ligand binding to receptors,
c) compounds which inhibit activation of intracellular signal pathways of the receptors,
d) compounds which inhibit or activate expression of a ligand or of a receptor of the VEGF or Tie receptor system,
e) delivery systems, such as antibodies, ligands, high-affinity binding oligonucleotides or oligopeptides, or liposomes, which target cytotoxic agents or coagulation-inducing agents to the endothelium via recognition of VEGF/VEGF receptor or Angiopoietin/Tie receptor systems,
f) delivery systems, such as antibodies, ligands, high-affinity binding oligonucleotides or oligopeptides, or liposomes, which are targeted to the endothelium and induce necrosis or apoptosis.

11. Pharmaceutical compositions according to claims 1-8 which comprise as compound II at least one of

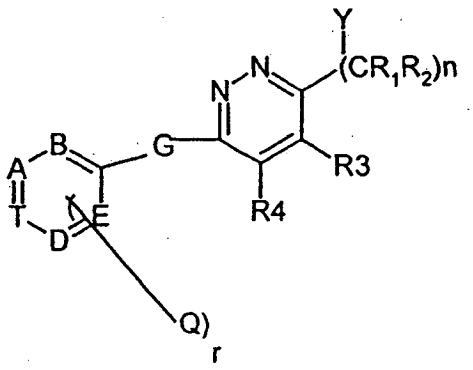
g) compounds which inhibit receptor tyrosine kinase activity,
h) compounds which inhibit ligand binding to receptors,
i) compounds which inhibit activation of intracellular signal pathways of the receptors,
j) compounds which inhibit or activate expression of a ligand or of a receptor of the VEGF or Tie receptor system,
k) delivery systems, such as antibodies, ligands, high-affinity binding oligonucleotides or oligopeptides, or liposomes, which target cytotoxic agents or coagulation-inducing agents to the endothelium via recognition of VEGF/VEGF receptor or Angiopoietin/Tie receptor systems,
l) delivery systems, such as antibodies, ligands, high-affinity binding oligonucleotides or oligopeptides, or liposomes, which are targeted to the endothelium and induce necrosis or apoptosis.

12. Pharmaceutical compositions according to claims 1-11 which comprise as compound I and/ or II at least one of Seq. ID Nos. 1-59.

13. Pharmaceutical compositions according to claims 1-11 which comprise as compound I and/ or II Seq. ID Nos. 34a

14. Pharmaceutical compositions according to claims 1-11 which comprise as compound I and/ or II at least one of sTie2, mAB 4301-42-35, scFv-tTF and/ or L19 scFv-tTFconjugate.

15. Pharmaceutical compositions according to claims 1-11 which comprise as compound I and/ or II at least one small molecule of genaral formula I

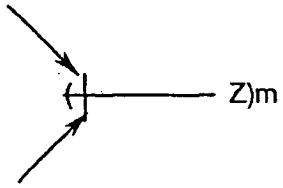


I,

in which

r has the meaning of 0 to 2,
n has the meaning of 0 to 2;

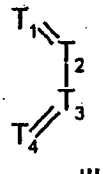
R_3 und R_4 a) each independently from eaxh other have the meaning of lower alkyl,
b) together form a bridge of general partial formula II,



II,

wherein the binding is via the two terminal C- atoms,
and

m has the meaning of 0 to 4; or
c) together form a bridge of partial formula III



III,

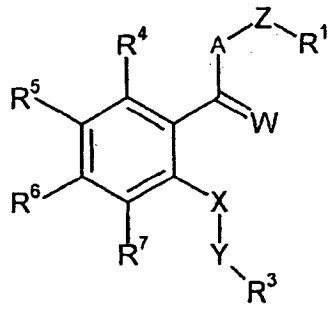
wherein one or two of the ring members T_1, T_2, T_3, T_4 has the meaning of nitrogen, and each others have the meaning of CH, and the bining is via the atoms T_1 and T_4;

G has the meaning of C_1-C_6 - alkyl, C_2 - C_6 - alkylene or C_2 - C_6 - alkenylene; or C_2 - C_6 - alkylene or C_3 -C_6-alkenylene, which are substituted with acyloxy or hydroxy; $-CH_2-O-$, $-CH_2-S-$, $-CH_2-NH-$, $-CH_2-O-CH_2-$, $-CH_2-S-CH_2-$, $-CH_2-NH-CH_2$, oxa (-O-), thia (-S-) or imino (-NH-),

A, B, D, E and T independently from each other have the meaning of N or CH , with the provisio that not more than three of these Substituents have the meaning of N,

Q	has the meaning of lower alkyl, lower alkyloxy or halogene,
R_1 and R_2	independently from each other have the meaning of H or lower alkyl,
X	has the meaning of imino, oxa or thia;
Y	has the meaning of hydrogene, unsubstituted or substituted aryl, heteroaryl, or unsubstituted or substituted cycloalkyl; and
Z	has the meaning of amino, mono- or disubstituted amino, halogen, alkyl, substituted alkyl, hydroxy, etherificated or esterificated hydroxy, nitro, cyano, carboxy, esterificated carboxy, alkanoyl, carbamoyl, N-mono- or N, N- disubstituted carbamoyl, amidino, guanidino, mercapto, sulfo, phenylthio, phenyl-lower-alkyl-thio, alkyl-phenyl-thio, phenylsulfinyl, phenyl-lower-alkyl-sulfinyl, alkylphenylsulfinyl, phenylsulfonyl, phenyl-lower-alkan-sulfonyl, or alkylphenylsulfonyl, whereas, if more than one rest Z is present (m≥2), the substituents Z are equal or different from each other, and wherein the bonds marked with an arrow are single or double bonds; or an N-oxide of said compound, wherein one ore more N-atoms carry an oxygene atom, or a salt thereof,

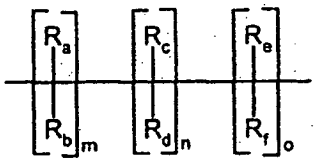
and/or a compound of genaral formula IV



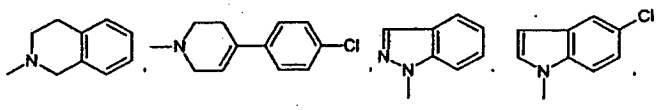
IV

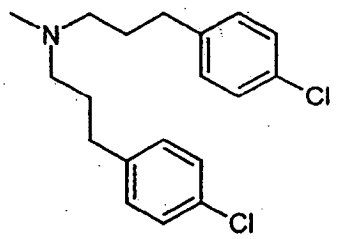
in which

A	has the meaning of group =NR^2,
W	has the meaning of oxygen, sulfur, two hydrogen atoms or the group =NR^8,
Z	has the meaning of the group =NR^{10} or =N-, -$N(R^{10})$-$(CH_2)_q$-, branched or unbranched C_{1-6}-Alkyl or is the group



or A, Z and R^1 together form the group





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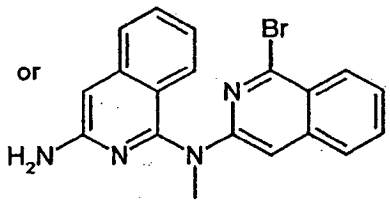
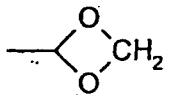


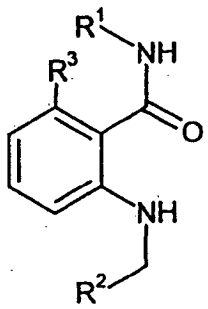
m, n and o	has the meaning of 0 - 3,
q	has the meaning of 1 - 6,
R_a, R_b, R_c, R_d, R_e, R_f	independently from each other have the meaning of hydrogen, C_{1-4} alkyl or the group $=NR^{10}$, and/ or R_a and/ or R_b together with R_c and or R_d or R_c together with R_e and/ or R_f form a bound, or up to two of the groups R_a-R_f form a bridge with each up to 3 C-atoms with R^1 or R^2,
X	has the meaning of group $=NR^9$ or =N-,
Y	has the meaning of group $-(CH_2)_p$,
P	has the meaning of integer 1-4,
R^1	has the meaning of unsubstituted or optionally substituted with one or more of halogene, C_{1-6}-alkyl, or C_{1-6}-alkyl or C_{1-6}-alkoxy, which is optionally substituted by one or more of halogen, or is unsubstituted or substituted aryl or heteroaryl,
R^2	has the meaning of hydrogen or C_{1-6}-alkyl, or form a bridge with up to 3 ring atoms with R_a-R_f together with Z or R_1,
R^3	has the meaning of monocyclic or bicyclic aryl or heteroaryl which is unsubstituted or optionally substituted with one or more of für halogen, C_{1-6}-alkyl, C_{1-6}-alkoxy or hydroxy,
R^4 ,R^5, R^6 and R^7	independently from each other have the meaning of hydrogen, halogene or C_{1-6}-alkoxy, C_{1-6}-alkyl or C_{1-6}-carboxyalkyl, which are unsubstituted or optionally substituted with one or more of halogene, or R^5 and R^6 together form the group



R^8, R^9 and R^{10}	independently from each other have the meaning of hydrogen or C_{1-6}-alkyl,

as well as their isomers and salts,

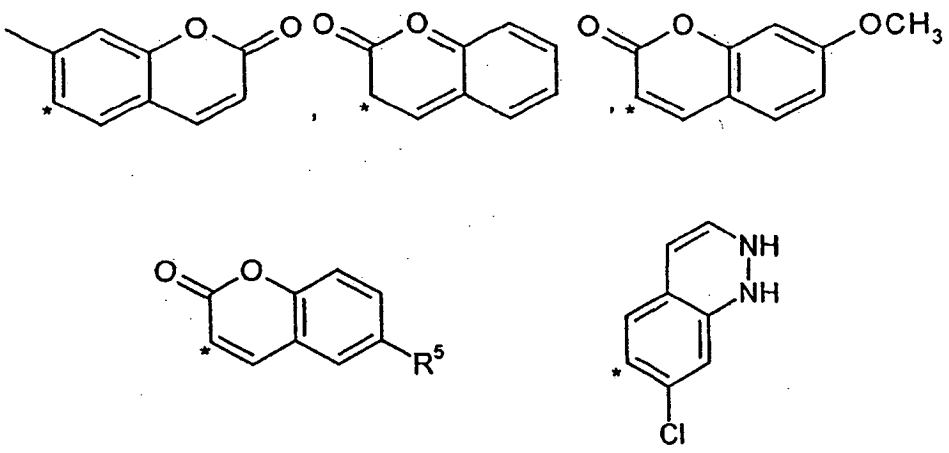
and/ or a compound of general formula V



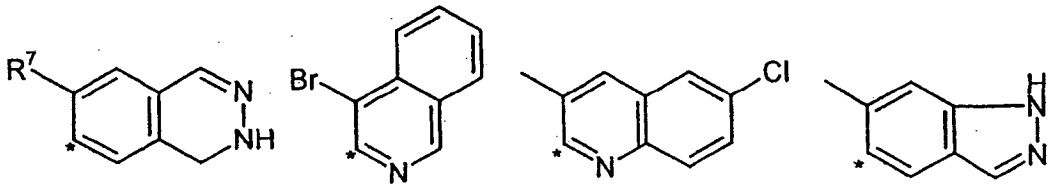
V,

in which

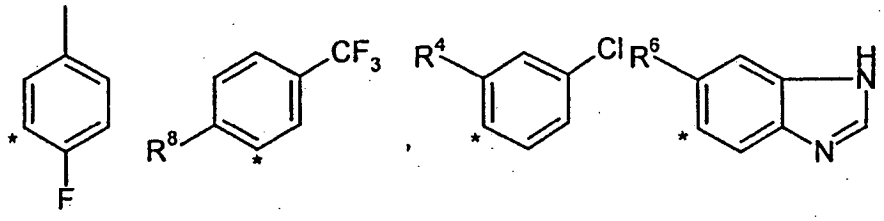
R^1 has the meaning of group



in which R^5 is chloro, bromo or the group $-OCH_3$,



in which R^7 is $-CH_3$ or chloro,

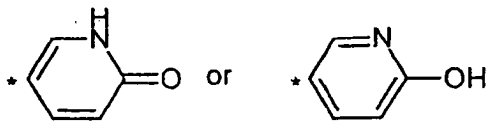


in which R^8 is $-CH_3$, fluoro, chloro or $-CF_3$

in which R^4 is fluoro, chloro, bromo, $-CF_3$, $-N{=}C$, $-CH_3$, $-OCF_3$ or $-CH_2OH$

in which R^6 is $-CH_3$ or chloro

R^2 has the meaning of pyridyl or the group



and

R^3 has the meaning of hydrogen or fluoro, as well as their isomers and salts.

16. Pharmaceutical compositions according to claim 15 which comprise as compound I and/ or II (4-Chlorophenyl)[4-(4-pyridylmethyl)-phthalazin-1-yl]ammonium hydrogen succinate

17. Pharmaceutical compositions according to claims 1-16 which comprise as compound I (4-Chlorophenyl)[4-(4-pyridylmethyl)-phthalazin-1-yl]ammonium hydrogen succinate, sTie2, mAB 4301-42-35, scFv-tTF and/ or L19 scFv-tTF conjugate, and as compound II (4-Chlorophenyl)[4-(4-pyridylmethyl)-phthalazin-1-yl]ammonium hydrogen succinatesTie2, mAB 4301-42-35, scFv-tTF and/ or L19 scFv-tTF conjugate, with the provisio that compound I is not identically to compound II.

18. Pharmaceutical compositions according to claims 1-17 which comprise as compound I (4-Chlorophenyl)[4-(4-pyridylmethyl)-phthalazin-1-yl]ammonium hydrogen succinate and as compound II sTie2, mAB 4301-42-35, scFv-tTF and/ or L19 scFv-tTF conjugate.

19. Pharmaceutical compositions according to claims 1-17 which comprise as compound I mAB 4301-42-35 and as compound II sTie2, and/ or scFv-tTF conjugate.

20. Pharmaceutical compositions according to claims 1-17 which comprise as compound I scFv-tTF conjugate and as compound II sTie2 and/ or mAB 4301-42-35.

21. Pharmaceutical compositions according to claims 1-17 which comprise as compound I L19 scFv-tTF conjugate and as compound II sTie2.

22. Use of pharmaceutical compositions according to claims 1-21, for the production of a medicament for the treatment of tumors, cancers, psoriasis, arthritis, such as rheumatoide arthritis, hemangioma, angiofribroma, eye diseases, such as diabetic retinopathy, neovascular glaukoma, kidney diseases, such as glomerulonephritis, diabetic nephropathie, maligneous nephrosclerosis, thrombic microangiopatic syndrome, transplantation rejections and glomerulopathy, fibrotic diseases, such as cirrhotic liver, mesangial cell proliferative diseases, artheriosclerosis, damage of nerve tissues, suppression of the ascites formation in patients and suppression of VEGF oedemas.

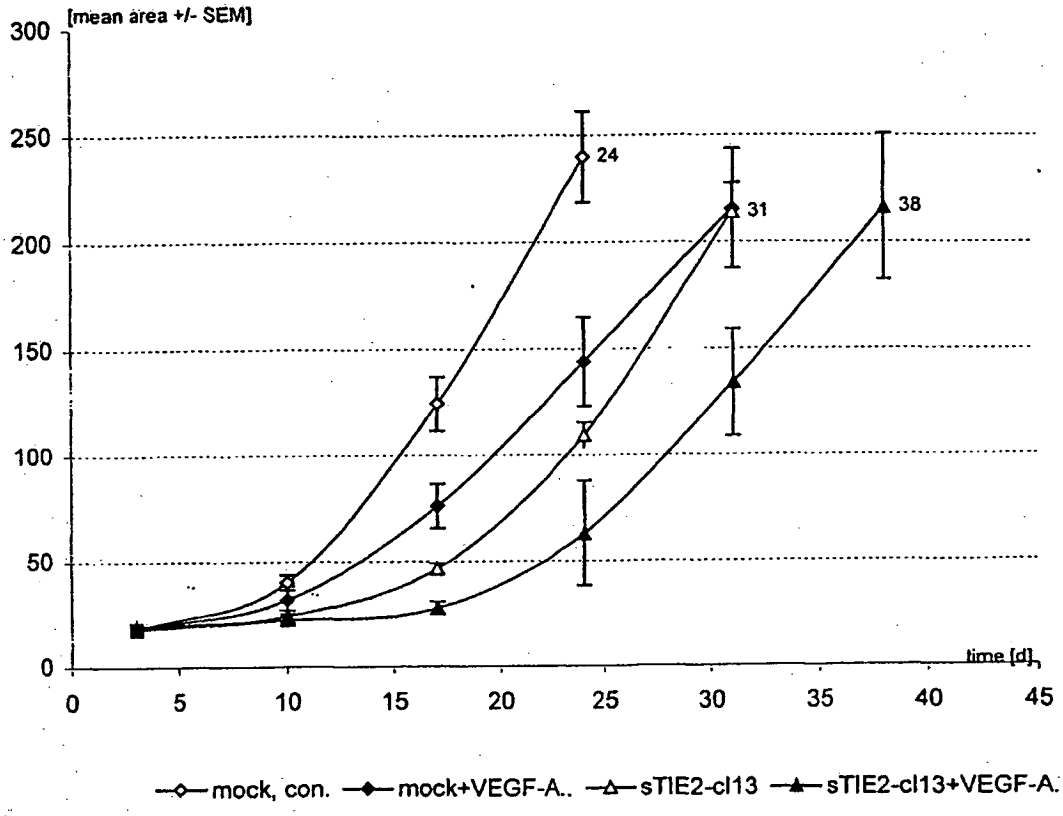

[mean area +/- SEM]
300
250
200
150
100
50
0
0
5
10
15
20
25
30
35
40
45
time [d]
24
31
38
mock, con.
mock+VEGF-A..
sTIE2-cl13
sTIE2-cl13+VEGF-A.

Fig. 1

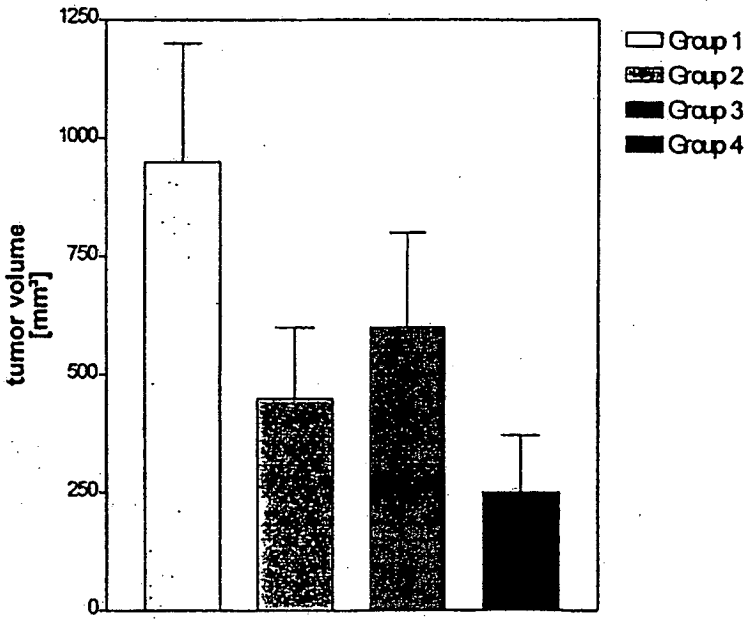

1250
1000
750
500
250
0
tumor volume [mm³]
Group 1
Group 2
Group 3
Group 4

Fig. 2

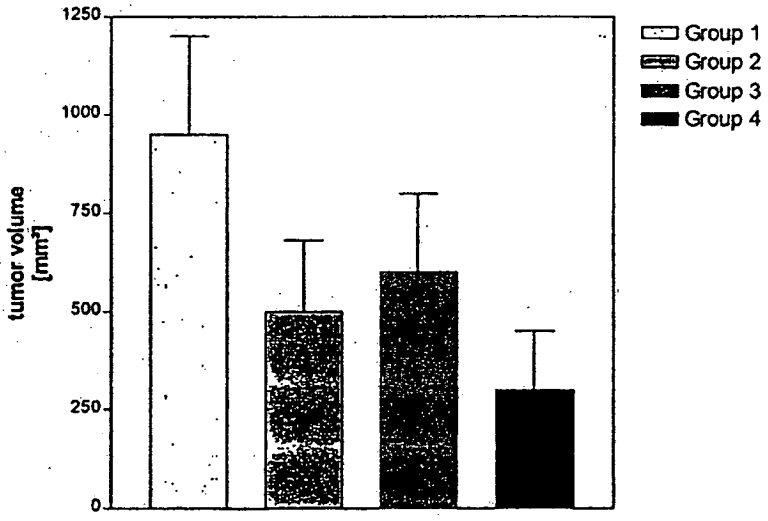

1250
1000
750
500
250
0
tumor volume [mm³]
Group 1
Group 2
Group 3
Group 4

Fig. 3

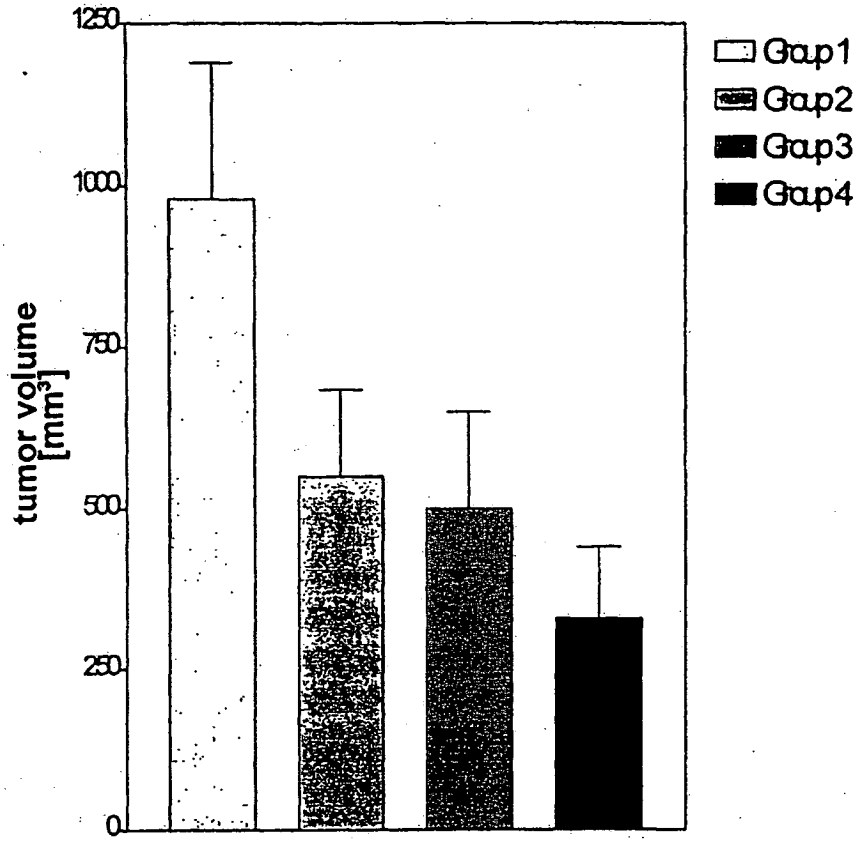


Fig. 4

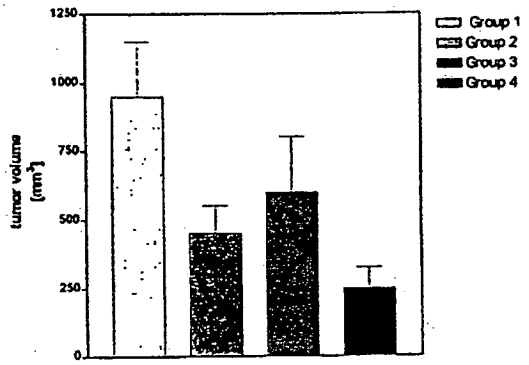

1250
1000
750
500
250
0
tumor volume [mm³]
Group 1
Group 2
Group 3
Group 4

Fig. 5

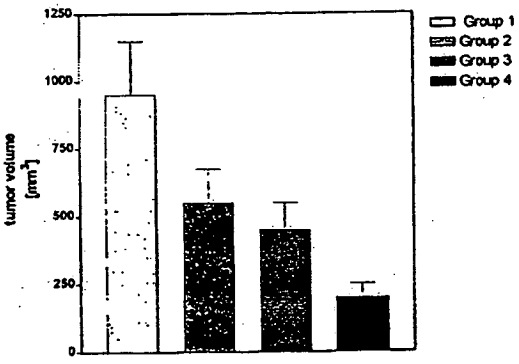

1250
1000
750
500
250
0
tumor volume [mm³]
Group 1
Group 2
Group 3
Group 4

Fig. 6

European Patent Office

PARTIAL EUROPEAN SEARCH REPORT

which under Rule 45 of the European Patent Convention shall be considered, for the purposes of subsequent proceedings, as the European search report

Application Number EP 00 25 0214

DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
X	WO 98 35958 A (NOVARTIS ERFINDUNGEN VERWALTUN ;CIBA GEIGY AG (CH); TRAXLER PETER) 20 August 1998 (1998-08-20) * abstract * * page 1, paragraph 1 *	1-15,21	A61K45/06
A	KOBLIZEK, THOMAS I. ET AL: "Angiopoietin-1 induces sprouting angiogenesis in vitro" CURR. BIOL. (1998), 8(9), 529-532 , XP000972727 * page 531, column 1, paragraph 2 *	1-21	
A	SCHLAEPPI, JEAN-MARC ET AL: "Targeting vascular endothelial growth factor (VEGF) for anti-tumor therapy, by anti-VEGF neutralizing monoclonal antibodies or by VEGF receptor tyrosine-kinase inhibitors." CANCER AND METASTASIS REVIEWS, (1999) VOL. 18, NO. 4, PP. 473-481. PRINT. , XP000972749 * page 477, column 2, paragraph 2 - page 478, column 1, paragraph 1 *	1-21	

TECHNICAL FIELDS SEARCHED (Int.Cl.7)

A61K

INCOMPLETE SEARCH

The Search Division considers that the present application, or one or more of its claims, does/do not comply with the EPC to such an extent that a meaningful search into the state of the art cannot be carried out, or can only be carried out partially, for these claims

Claims searched completely

Claims searched incompletely

Claims not searched

Reason for the limitation of the search:

see sheet C

Place of search	Date of completion of the search	Examiner
THE HAGUE	12 January 2001	Leherte, C

CATEGORY OF CITED DOCUMENTS

X : particularly relevant if taken alone
Y : particularly relevant if combined with another document of the same category
A : technological background
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D : document cited in the application
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EPO FORM 1503 03.82 (P04C07)

European Patent Office

INCOMPLETE SEARCH
SHEET C

Application Number
EP 00 25 0214

Claim(s) searched completely:
17-21

Claim(s) searched incompletely:
1-16, 22

Reason for the limitation of the search:

Present claims 1-16, 22 relate to compounds defined by reference to desirable characteristics or properties, namely :
" agents which modulates the biological function of one or several of the VEGF/VEGF receptor systems",
" agents which modulate the biological function of one or several of the Angiopoietin/Tie receptor systems",
" agents which are targeted to the endothelium via one or several of the VEGF/VEGF receptor systems",
" agents which are targeted to the endothelium",
" agents which are targeted to the endothelium via one or several of the Angiopoietin/Tie receptor systems",
" agents which interfere with both the function of one or several of the VEGF/VEGF receptor systems and the function of the Angiopoietin/Tie receptor systems",
" compounds which inhibit receptor tyrosine kinase activity",
" compounds which inhibit ligand binding to receptors",
" compounds which inhibit activation of intracellular signal pathways of the receptors",
" compounds which inhibit or activate expression of a ligand or of a receptor of the VEGF or Tie receptor systems",
" delivery systems, which target cytotoxic agents or coagulation-inducing agents to the endothelium",
"delivery systems, such as antibodies, ligands, high-affinity binding oligonucleotides or oligopeptides, or liposomes, which are targeted to the endothelium and induce necrosis or apoptosis",
" compounds of general formula I, IV or V".
The claims cover all compounds having those characteristics, properties or these general formulas, whereas the application provides support within the meaning of Article 84 EPC and/or disclosure within the meaning of Article 83 EPC for only a very limited number of such compounds. In the present case, the claims so lack support, and the application so lacks disclosure, that a meaningful search over the whole of the claimed scope is impossible. Independent of the above reasoning, the claims also lack clarity (Article 84 EPC). An attempt is made to define the compounds by reference to a result to be achieved. Again, this lack of clarity in the present case is such as to render a meaningful search over the whole of the claimed scope impossible. Consequently, the search has been carried out for those parts of the claims which appear to be clear, supported and disclosed, namely those parts relating to the pharmaceutical compositions mentioned in the claims 17-21, with due regard to the therapeutic applications mentioned in claim 22.

ANNEX TO THE EUROPEAN SEARCH REPORT ON EUROPEAN PATENT APPLICATION NO. EP 00 25 0214

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12-01-2001

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9835958 A	20-08-1998	AU 6621898 A	08-09-1998
		BR 9807685 A	21-03-2000
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		CZ 9902853 A	17-11-1999
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		SK 109699 A	13-03-2000
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